What is claimed is:

- 1. An expandable intraluminal stent comprising a main body portion having a first end portion, a second end portion, a middle portion and a flow passage defined therethrough, at least a portion of the first end portion having a biocompatible coating.
- 2. The stent of claim 1 wherein the coating comprises a biocompatible polymer.
- The stent of claim 1 wherein the coating comprises a biodegradable polymer.
- 10 4. The stent of claim 1 wherein the first end portion comprises an inner surface, an outer surface, an end and an edge, the coating covering the end and at least a portion of the edge of the first end portion.
 - 5. The stent of claim 1 wherein at least a portion of the second end portion has a biocompatible coating.
- 15 6. The stent of claim 5 wherein the second end portion comprises an outer surface, an inner surface, an end, and an edge, the coating covering the end and at least a portion of the edge of the second end portion.
 - 7. The stent of claim 5 wherein the coating comprises a biocompatible polymer.
- 20 8. The stent of claim 1 wherein the coating includes a drug.
 - 9. The stent of claim 8 wherein the drug comprises TAXOL.
 - 10. The stent of claim 1 wherein the coating includes a bioadhesive.
 - 11. The stent of claim 1 wherein the coating comprises a plurality of layers.
- 12. The stent of claim 11 wherein the plurality of layers is comprised of the same coating material.
 - 13. The stent of claim 11 wherein the plurality of layers is comprised of different coating materials.
 - 14. The stent of claim 12 wherein at least one of the layers includes a drug.
 - 15. The stent of claim 5 wherein the coating comprises a plurality of layers.
- The stent of claim 15 wherein the plurality of layers is comprised of the same coating material.

- 17. The stent of claim 15 wherein the plurality of layers is comprised of different materials.
 - 18. The stent of claim 15 wherein at least one of the layers includes a drug.
- 19. An expandable intraluminal stent comprising a main body portion having a first end portion, a second end portion, a middle portion and a flow passage
- defined therethrough, and a sleeve of biocompatible material connected to the first end portion.
- 20. The stent of claim 19 wherein the sleeve comprises a biocompatible polymer.
- The stent of claim 19 wherein the coating comprises a biodegradable polymer.
 - 22. The stent of claim 19 wherein the sleeve includes apertures.
 - 23. The stent of claim 19 wherein the sleeve includes a drug.
 - 24. The stent of claim 23 wherein the drug comprises TAXOL.
- 15 25. The stent of claim 19 wherein the sleeve comprises a plurality of layers.
 - 26. The stent of claim 25 wherein the plurality of layers is comprised of the same coating material.
- 27. The stent of claim 25 wherein the plurality of layers is comprised of 20 different materials.
 - 28. The stent of claim 25 wherein at least one of the layers includes a drug.
 - 29. The stent of claim 19 further comprising a second sleeve connected to the second end portion.
- 30. The stent of claim 29 wherein the sleeve comprises a biocompatible polymer.
 - 31. The stent of claim 29 wherein the sleeve includes apertures.
 - 32. The stent of claim 29 wherein the sleeve includes a drug.
 - 33. The stent of claim 29 wherein the sleeve comprises a plurality of layers.
- 30 34. The stent of claim 33 wherein the plurality of layers is comprised of the same coating material.

- 35. The stent of claim 33 wherein the plurality of layers is comprised of different materials.
 - 36. The stent of claim 33 wherein at least one of the layers includes a drug.
- 37. An expandable intraluminal stent comprising a main body portion having a first end portion, a second end portion, a middle portion and a flow passage defined therethrough, the first end portion being polished to provide a smooth first end portion.
 - 38. The stent of claim 37 further comprising a polished second end portion to provide a smooth second end portion.
- 10 39. An expandable intraluminal stent comprising a main body portion having a first end portion, a second end portion, a middle portion and a flow passage defined therethrough, the first end portion being heat treated to provide flexibility.
 - 40. The stent of claim 39 further comprising a heat treated second end portion to provide flexibility.
- 15 41. An expandable intraluminal stent comprising a main body portion having a first end portion, a second end portion, a middle portion and a flow passage defined therethrough, the first end portion constructed in a manner so as to be more flexible than the middle portion.
 - 42. The stent of claim 41 wherein the stent comprises a looser mesh first end portion than the middle portion.
 - 43. The stent of claim 41 further comprising the second end portion constructed in a manner so as to be more flexible than the middle portion.
 - 44. The stent of claim 43 wherein the stent comprises a looser mesh second end portion than the middle portion.
- 25 45. The stent of claim 1 wherein the coating comprises an RGD peptidecontaining compound.
 - 46. The stent of claim 8 wherein the drug comprises 5-flroracil.
 - 47. The stent of claim 8 wherein the drug comprises Tranilast.
 - 48. The stent of claim 8 wherein the drug comprises Tropidil.
- The stent of claim 8 wehrien the drug comprises Probucol.

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